Section: 21.0 Hepatitis B, acute/Delta Hepatitis

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Hepatitis B, acute

Overview^(1,2)

For a complete description of hepatitis B, refer to the following texts:

- Control of Communicable Diseases Manual (CCDM).
- Red Book, Report of the Committee on Infectious Diseases.

Case Definition(3)

Clinical Definition

Hepatitis B:

An acute illness with:

- a) Discrete onset of symptoms
- b) Jaundice or elevated serum aminotransferase levels

Perinatal Hepatitis B:

Perinatal hepatitis B in the newborn may range from asymptomatic to fulminant hepatitis.

Laboratory criteria for diagnosis

Hepatitis B:

- 1. IgM antibody to hepatitis B core antigen (anti-HBc IgM) positive (if done) or hepatitis B surface antigen (HBsAg) positive
- 2. IgM anti-HAV negative (if done)

Hepatitis B Perinatal:

Hepatitis B surface antigen (HbsAg) positive

Case classification

Hepatitis B:

Confirmed: a case that meets the clinical case definition and is laboratory confirmed.

Case Definitions for Probable, Suspect, Old and Carrier Cases: (5)

Clinical description:

Cases in these categories may or may not have an acute onset of illness or symptoms and may or may not have elevated liver enzyme test results. Additional information is required to correctly classify the case.

Probable: A case that is positive for Hepatitis B core IgM Antibody (Anti-HBc IgM) but symptom history is currently unavailable and the case has not been reported previously. This case should remain open until information is available to reclassify the case. ⁽⁵⁾

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Suspect: A case that is positive for Hepatitis B Surface Antigen (HBsAg) and does not have symptoms and is either negative for Hepatitis B Core IgM (Anti-HBc IgM) or Anti-HBc IgM results are not available. (5)

Old: A case that is negative for Hepatitis B Surface Antigen (HBsAg) and does not have symptoms and is positive for Hepatitis B Core Antibody Total (Anti-HBc) and is negative for Hepatitis B Core IgM (Anti-HBc IgM) or Anti-HBc Igm results are not available. (5)

Carrier: A case that has been positive for Hepatitis B Surface Antigen (HBsAg) for more than six months OR a case that is positive for Hepatitis B Surface Antigen (HBsAg) and is positive for Hepatitis B Core Antibody Total (Anti-HBc) and is negative for Hepatitis B Core Igm (Anti-HBc IgM). (5)

Case Definition	Discrete onset of symptoms	Jaundice or elevated liver functions	HBsAg	Anti-HBs	Anti-HBc	Anti-HBc IgM	HAVIgm
Confirmed	Yes	Yes	+	Unk	Unk	+ (If done)	- (If done)
Probable	Unk	Unk	+	Unk	Unk	+	Unk
Suspect	No	No	+	Unk	Unk	- (If done)	Unk
Old	No	No	-	Unk	+	-	Unk
Carrier*	No	No	+	Unk	+	-	Unk

^{*} A case that has been positive for HBsAg for more than six months also fits within the case definition of carrier.

Perinatal Hepatitis B:

Confirmed: HBsAg positivity in any infant aged >1-24 months who was born in the United States or in U.S territories to an HBsAg-positive mother.

Comment:

1. Infants born to HBsAg-positive mothers should receive hepatitis B immune globulin (HBIG) and the first dose of hepatitis B vaccine within 24 hours of birth, followed by the second and third doses of vaccine at 1 and 6 months of age, respectively. Post-vaccination testing for HBsAg and the antibody to HBsAg (Anti-HBsAg) is recommended from 3 to 6 months following completion of the vaccine series. If HBIG and the initial dose of vaccine are delayed for >1 month after birth, testing for HBsAg may determine if the infant is already infected. If vaccine is not available at birth, the first dose should be given prior to discharge, but not later than 7 days after birth. (4)

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<u>Information Needed for Investigation</u>

Verify the diagnosis. What laboratory tests were conducted? What were the results? What are the case's clinical symptoms? Is this an acute case or is this a hepatitis B carrier? Do a search in MOHSIS to reveal any previous history of hepatitis B. What is the individual's vaccination status?

Contact the District Immunization Representative immediately if the case is prenatal or perinatal regardless of case classification.

Establish the extent of illness. Determine if household or other close contacts are, or have been ill, by contacting the health care provider, patient or family member. **Contact the District Communicable Disease Coordinator** if cases are in high-risk settings or jobs such as health care.

Case/Contact Follow-Up And Control Measures

Determine the source of infection:

- Is this a perinatal exposure?
- Did this case have a percutaneous or permucosal exposure?
- Did this case have unprotected sex with a hepatitis B positive partner?
- Have there been other cases linked by time, place or person?
- Did this person have household exposure to another hepatitis B case or carrier?

Determine the risk of infection to others and to contact those who may be at risk:

- Is the case pregnant?
- Does this case have unprotected sex with multiple partners?
- Does this case participate in the use of intravenous drugs or any activity that may involve the sharing or re-use of needles such as tattooing/body piercing?

Control Measures

See the Viral Hepatitis B and Delta Hepatitis sections of the <u>Control of Communicable Diseases Manual (CCDM)</u>, "Control of Patient, Contacts and the Immediate Environment".

See the Hepatitis B and Hepatitis D sections of the Red Book.

Contact the District Immunization Representative to arrange for vaccination of household contacts and other high-risk individuals. Further information on who is eligible for these vaccinations can be found in the Section of Vaccine Preventable and Tuberculosis Disease Elimination (SVPTDE) Immunization Guidelines Manual.

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Laboratory Procedures

- 1. For confirmation of a suspected case of hepatitis B, use a Missouri State Public Health Laboratory (SPHL) virus serology kit.
- 2. The SPHL does <u>NOT</u> perform Delta hepatitis testing. If Delta hepatitis is suspected, recommend that the physician submit the specimen to a commercial laboratory for testing. If a commercial laboratory is not an option, contact the District Communicable Disease Coordinator and he/she will coordinate sending the specimen to CDC.
- 3. Draw one red-top tube of blood, using standard precautions.
- 4. Either send serum OR whole blood:

To send serum:

- Allow blood to stand for 30 minutes to 1 hour
- Centrifuge at 2000 rpm for 5 minutes
- Pour clear serum into another red top tube*
- Discard clot tube safely

NOTE: Hot or cold weather conditions can hemolyze whole blood, making it unsuitable for testing. Specimens should be sent as serum under these conditions.

- 5. Wipe off outside of tube with alcohol.
- 6. Label tube carefully; include patient's name and date sample obtained. Samples without this information on both the tube and request form *will not be tested and will be discarded*.
- 7. Fill out lab form "Hepatitis Test Request", as completely as you can. If you are screening contacts mark the box labeled "Hepatitis B-EIA for (HBsAg)".
- 8. Immediately mail blood or serum; include form.
- 9. If the test is requested by a local public health agency, state the name of the LPHA on the lab form. If submitted by a private physician, he/she will be charged.

*The state lab cannot provide the extra red top tubes.
FOR ADDITIONAL INFORMATION, CONTACT THE DISTRICT
COMMUNICABLE DISEASE COORDINATOR

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Interpretation of Specific Serological Tests for Hepatitis B:

Antigens: Parts of the virus itself.

Test	Interpretation
	<u>Interpretation</u>
HBsAg	Infectious – Surface antigen appears in the serum as early
Hepatitis B Surface Antigen	as 1-2 and as late as 11-12 weeks after exposure to HBV.
	It is usually the first viral marker to appear in the blood
	after HBV infection. In self-limited infections (not a
	carrier), HBsAg remains detectable in the blood for 1-6
	weeks although it may persist for as long as 20 weeks.
	95% of patients are HBsAg positive at the onset of
	symptoms and jaundice. HBsAg generally disappears
	with recovery. In some patients, HBsAg clears rapidly
	and may be absent at the time the patient is tested.
	Between 5-10% of patients infected with HBV do not
	clear HBsAg and become carriers. A patient with HBsAg
	that persists beyond 6 months after acute infection or a
	previously positive HBsAg test is considered a carrier.
	HBsAg persists indefinitely in carriers. Look at antibody
	tests and any previous viral hepatitis serology tests to
	determine whether the case has been recently infected or is
	a carrier.
HBeAg	Highly Infectious- HBeAg is another regular and early
Hepatitis B "Early" Antigen	marker of HBV infection. HBeAg appears simultaneously
	or within a few days of the appearance of HBsAg in most
	primary infections. Its titer peaks and then declines in
	parallel with HBsAg. HBeAg usually disappears just
	before the disappearance of HBsAg in self-limited
	infections. Patients who remain HBeAg positive for 10
	weeks or longer appear likely to become carriers. Look at
	antibody tests and any previous viral hepatitis serology
	tests to determine whether the case has been recently
	infected or is a carrier.

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Antibodies: Proteins developed by the body's immune system in response to antigens.

<u>Test</u>	<u>Interpretation</u>
Anti-HBc	<u>Indeterminate infectiousness</u> - Anti-HBc titers usually rise
Hepatitis B Core Antibody	during the period of HBsAg positivity, level off, and
Total	slowly fall after HBsAg becomes undetectable. Anti-HBc
	can be detected for 5 to 6 years after acute infection in
	both carriers and non-carriers. Anti-HBc is the
	combination of IgM and IgG class antibodies (Anti-HBc
	IgM and Anti-HBc IgG). Look at Anti-HBc Igm to
	determine recent infection. Look at antigen tests to
	determine whether the case is currently infectious.
Anti-HBc Igm or HBcIgm	<u>Indeterminate infectiousness, Recent infection</u> -Anti-HBc
Hepatitis B Core IgM	IgM is one class of Hepatitis B Core antibodies. Anti-
Antibody	HBc has been found in almost all patients with acute
	hepatitis B. Anti-HBc Igm rapidly decreases in titer after
	infection and is no longer detectable within 6-24 months.
	A positive Anti-HBc Igm serology is an indication of
	recent infection. A chronic hepatitis B carrier should test
	negative for Anti-HBc Igm. Look at antigen tests to
	determine whether the case is currently infectious.
Anti-HBc IgG or HBcIgG	<u>Indeterminate infectiousness</u> -The individual was infected
Hepatitis B Core IgG	at some time in the past. This test alone does not give any
Antibody	indication of when in the past. Look at Anti-HBc Igm to
	determine recent infection. Look at antigen tests to
	determine whether the case is currently infectious.
Anti-HBe	Probably less infectious- Anti-HBe appears in most
Hepatitis B "Early"	patients at the time HBeAg becomes undetectable or
Antibody	shortly thereafter. Anti-HBe persists for 1 to 2 years after
	resolution of HBV infection. Look at Anti-HBc Igm to
	determine recent infection. Look at antigen tests to
A CHE HE AL	determine whether the case is currently infectious.
Anti-HBs or HBsAb	Immune- Anti-HBs develops after a resolved infection and
Antibody to Hepatitis B	is a marker of long-term immunity. This is the only
Surface Antigen	positive test for an uninfected, vaccinated individual. It
	may take several months for this antibody to appear after
	infection or vaccination and in some cases does not appear
	at all. However, this antibody will likely persist for
	several years when it does appear.

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Interpretation of Several Combinations of Hepatitis B Serologic Tests

	Test	s for HBV		<u>Interpretation</u>
HBsAg	Anti-HBs	Anti-HBc	Anti-HBc IgM	
+	_	_	_	Very recent, acute HBV infection (infectious)
+	_	+	+	Acute HBV infection (infectious)
+	_	+	_	Chronic HBV infection with HBsAg carriage (infectious)
_	_	+	+	Acute HBV infection, anti—HBs has not yet appeared (may be infectious)
_	+	+	+	Resolving acute HBV infection (may be infectious)
_	+	+	_	HBV infection in the remote past (Immune)
_	_	+	_	HBV infection in the remote past (Immune)
_	_	_	Not Tested	No hepatitis B infection, if liver abnormalities exist they are due to another virus, toxin or condition
_	+	_	Not Tested	Post hepatitis B vaccine or post HBIG

Reporting Requirements

Hepatitis B is a Category II disease and shall be reported to the local health authority or to the Missouri Department of Health and Senior Services (DHSS) within 3 days of first knowledge or suspicion.

- 1. For all cases, complete a Disease Case Report form (CD-1).
- 2. For confirmed, probable, suspect and perinatal cases, complete the "Viral Hepatitis Case Report" form (CDC 53.1), front and back. Detach and complete the worksheet on the back (this is necessary to prevent the carbons from marking on the front of the form). Staple the completed worksheet to the form.
- 3. If this is a prenatal or perinatal case, contact the District Immunization Representative. Fill out the "Prenatal Hepatitis B Case Report" form (IMMP-29) and the "Perinatal Hepatitis B Case Report" form (IMMP-29A). These forms, policy, and guidelines can be found in the Section of Vaccine Preventable and Tuberculosis Disease Elimination (SVPTDE) Immunization Guidelines Manual.
- 4. If prophylaxis is provided using hepatitis B vaccine and/or HBIG supplied by DHSS, SVPTDE, complete forms and follow protocol in the Immunization Guidelines Manual.
- 5. Entry of the complete CD-1 into the MOHSIS database negates the need for the paper CD-1 to be forwarded to the District Health Office.
- 6. Send the completed secondary investigation form(s) to the District Health Office.
- 7. All outbreaks or "suspected" outbreaks must be reported as soon as possible (by phone, fax or e-mail) to the District Communicable Disease Coordinator. This can be accomplished by completing the Missouri Outbreak Surveillance Report (CD-51).
- 8. Within 90 days from the conclusion of an outbreak, submit the final outbreak report to the District Communicable Disease Coordinator.

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References

- 1. Chin, James, ed. "Hepatitis B and Delta Hepatitis." <u>Control of Communicable Diseases Manual</u>, 17th Ed. Washington, D.C.: American Public Health Association., 2000: 243-251, 253-255.
- 2. American Academy of Pediatrics. "Hepatitis B and Hepatitis D." In: Pickering, LK, ed. 2000 Red Book: Report of the Committee on Infectious Diseases. 25th ed. Elk Grove Village, IL. 2000: 289-302, 306-308.
- Centers for Disease Control and Prevention. <u>Case Definitions for Infectious</u> <u>Conditions Under Public Health Surveillance</u>. MMWR 1997:46 (No.RR-10): 18-19.
- **4.** Atkinson, William, et al. Eds. <u>Epidemiology and Prevention of Vaccine-Preventable Diseases</u>; 7th Ed.: Atlanta: National Immunization Program, Centers for Disease control and Prevention, 2002: 223-245.
- **5.** Missouri Department of Health and Senior Services Section of Communicable Disease Control and Veterinary Public Health surveillance case definition.
- **6.** Missouri Department of Health and Senior Services <u>Section of Vaccine-Preventable and Tuberculosis Disease Elimination Immunization Guidelines</u>, Section 345, Subsection 345.07.
- 7. Mandell, GL, Bennett, JE, and Dolin, R, ed. *Mandell Douglas and Bennett's Principles and Practice of Infectious Diseases*, 5th ed. New York: Churchill Livingstone, 2000.

Other Sources of Information

Evans, Alfred S. and Richard A. Kaslow, Eds. <u>Viral Infections of Humans Epidemiology</u> and <u>Control</u>; 4th ed. Eds.. New York: Plenum, 1997: 375-387.

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Delta Hepatitis

$Overview^{(1,2)}$

For a complete description of Delta hepatitis, refer to the Delta hepatitis section in the following texts:

- Control of Communicable Diseases Manual (CCDM).
- Red Book, Report of the Committee on Infectious Diseases.

Case Definition(3)

Clinical case definition

An acute illness with:

- a) Discrete onset of symptoms
- b) Jaundice or elevated serum aminotransferase levels

Laboratory criteria for diagnosis

Delta hepatitis: HBsAg or IgM anti-HBc positive and antibody to hepatitis delta virus positive

Case classification:

Confirmed: a case that meets the clinical case definition and is laboratory confirmed.

*Delta hepatitis is not a nationally notifiable disease.

<u>Information Needed for Investigation</u>

Verify the diagnosis. What laboratory tests were conducted? What were the results? What are the case's clinical symptoms? Is this person documented as an acute case of hepatitis B or hepatitis B carrier? (Delta hepatitis is only present in individuals with hepatitis B).

Establish the extent of illness. Determine if household or other close contacts are, or have been ill, by contacting the health care provider, patient or family member.

Contact the District Communicable Disease Coordinator. Due to the low prevalence of hepatitis D, one occurrence of disease is significant.

Case/Contact Follow-Up And Control Measures

Determine the source of infection:

- Is this a perinatal exposure?
- Did this case have a percutaneous or permucosal exposure?
- Did this case have unprotected sex with a delta hepatitis positive partner?
- Have there been other cases linked by time, place or person?
- Did this person have household exposure to another delta hepatitis case?

Determine the risk of infection to others and to contact those who may be at risk:

- Does this case have unprotected sex with multiple partners?
- Does this case participate in intravenous use of illegal drugs?

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Control Measures

See the Delta Hepatitis section of the <u>Control of Communicable Diseases Manual</u> (CCDM), "Control of patient, contacts and the immediate environment".

See the Hepatitis D section of the <u>Red Book</u>.

Laboratory Procedures

The SPHL does <u>NOT</u> perform Delta hepatitis testing. If Delta hepatitis is suspected, recommend that the physician submit the specimen to a commercial laboratory for testing. If submitting the specimen to a commercial laboratory is not an option, contact the District Communicable Disease Coordinator and he/she will coordinate sending the specimen to CDC.

FOR ADDITIONAL INFORMATION, CONTACT YOUR DISTRICT COMMUNICABLE DISEASE COORDINATOR.

Reporting Requirements

Delta Hepatitis is a Category II disease and shall be reported to the local health authority or to the Missouri Department of Health and Senior Services (DHSS) within 3 days of first knowledge or suspicion.

- 1. For all cases complete a Disease Case Report form (CD-1) and the "Viral Hepatitis Case Report" form (CDC 53.1), front and back. Detach and complete the worksheet on the back (this is necessary to prevent the carbons from marking on the front of the form). Staple the completed worksheet to the form.
- 2. Entry of the complete CD-1 into the MOHSIS database negates the need for the paper CD-1 to be forwarded to the District Health Office.
- 3. Send the completed secondary investigation form to the District Health Office.
- 4. All outbreaks or "suspected" outbreaks must be reported as soon as possible (by phone, fax or e-mail) to the District Communicable Disease Coordinator. This can be accomplished by completing the Missouri Outbreak Surveillance Report (CD-51).
- 5. Within 90 days from the conclusion of an outbreak, submit the final outbreak report to the District Communicable Disease Coordinator.

References

- 1. Chin, James, ed. "Delta Hepatitis." <u>Control of Communicable Diseases Manual</u>, 17th ed. Washington, D.C.: American Public Health Association, 2000: 253-255.
- 2. American Academy of Pediatrics. "Hepatitis D." In: Pickering, LK, ed. 2000 Red Book: Report of the Committee on Infectious Diseases. 25th ed. Elk Grove Village, IL. 2000: 306-308.
- 3. Centers for Disease Control and Prevention. <u>Case Definitions for Infectious Conditions Under Public Health Surveillance</u>. MMWR 1997:46 (No.RR-10).
- 4. Missouri Department of Health and Senior Services Section of Communicable Disease Control and Veterinary Public Health surveillance case definition.

Other Sources of Information

- 1. Mandell, GL, Bennett, JE, and Dolin, R, ed. *Mandell Douglas and Bennett's Principles and Practice of Infectious Diseases*, 5th ed. New York: Churchill Livingstone, 2000.
- 2. Evans, Alfred S. and Richard A. Kaslow, Eds. <u>Viral Infections of Humans Epidemiology and Control</u>; 4th ed. Eds. New York: Plenum, 1997: 375-387.

Hepatitis B

Fact Sheet

What is hepatitis B?

Hepatitis B (HBV) formerly known, as serum hepatitis is a liver disease caused by the hepatitis B virus. An estimated 140,000 Americans are infected each year. Approximately 1-1.25 million people are chronically infected and are considered to be carriers of the virus.

Who gets hepatitis B?

Anyone can get hepatitis B, but those at greatest risk include:

- Babies born to mothers who are infectious with the hepatitis B virus (including carriers)
- Injection drug users (or persons who share needles);
- Persons in occupations where there is contact with infected blood and other infectious body fluids which includes saliva - (persons who work as health care workers, first responders, etc.):
- Sexually active homosexual and bisexual men;
- Anyone with multiple sex partners;
- Men and women who have a recent diagnosis of another sexually transmitted disease
- Hemodialysis patients;
- Persons who receive unscreened blood products;
- Certain household contacts of an infected person;
- Household contacts and sexual partners of hepatitis B carriers
- Inmates of long-term correctional facilities
- Persons from countries where hepatitis B is widespread (certain areas of Asia and Africa);
- Persons in the United States who are Alaskan Natives or Pacific Islanders.
- Persons receiving tattoos/body piercings.

How is the virus spread?

Hepatitis B virus can be found in the blood of infected persons; it is also present in lesser amounts in semen, vaginal secretions, amniotic fluid, body fluids containing blood, unfixed tissues and organs, and saliva. Hepatitis B virus is spread through sexual contact (homosexual and heterosexual), needle sharing, needle stick injury, mucous membrane exposure, and direct contact with infected body fluids. Hepatitis B virus is not spread by casual contact.

What are the symptoms of hepatitis B?

Many persons with hepatitis B infection have no symptoms at all, or they may be mild and flulike (i.e.: loss of appetite, nausea, vomiting, diarrhea, fatigue). Some patients may notice darkened urine (iced-tea colored), light clay colored stools or yellowing of the skin and eyes (jaundice). A few persons will experience a more severe disease syndrome and may die of overwhelming liver failure.

How soon do symptoms appear?

The symptoms, if they appear, may occur as early as 45 days to as long as 180 days following exposure, the average is 60-90 days. The variation in time for onset of symptoms is related in part to the amount of virus transmitted at the time of exposure.

How is hepatitis B diagnosed?

Since there are several types of viral hepatitis, a blood test is needed to determine which type of hepatitis is present.

For how long is a person able to spread the virus?

The virus can be present in blood and other body fluids several weeks before symptoms appear and generally for several months afterward. About 1 in 10 adults and 3 of 10 children with hepatitis B will go on to become ongoing carriers of the virus for the rest of their lives even when they do not have symptoms. As long as the virus is present, it can be spread to others. Persons with hepatitis B infection should not donate blood, organs or tissues.

What is the treatment for hepatitis B?

There are no special medicines to treat a person with acute hepatitis B disease. Generally, rest and supportive treatment is all that is needed. Carriers of hepatitis B infection may benefit from treatment with a medication called interferon.

What precautions should hepatitis B carriers take?

Hepatitis B carriers should practice good hygiene so that close contacts are not directly exposed to their blood or other body fluids. Carriers must not share razors, toothbrushes or any other object that may become contaminated with blood. In addition, household members, particularly sexual partners, should be immunized with hepatitis B vaccine. Carriers should practice "safe sex"; use barrier precautions, if their partners are not immunized. It is important for carriers to inform their dentists and other health care providers of their carrier state.

Can hepatitis B be prevented?

Yes, a vaccine to prevent hepatitis B has been available since 1982. It is safe, effective and recommended for all newborns, 11-12 year olds and for any person at risk for getting infected (see page 1, "Who gets hepatitis B?"). A special hepatitis B immune globulin (HBIG) is also available for people who are not vaccinated and are exposed to the virus. If you are exposed to hepatitis B, consult a doctor or your local health department immediately.

Missouri Department of Health and Senior Services Section of Communicable Disease Control and Veterinary Public Health Phone: (800) 392-0272 This questionnaire is authorized by law (Public Health Service Act, 42 USC 2k1). Atthough response to the questions is voluntary, cooperation of the patient is necessary for the study and control of the decision of the patient is necessary for the study and control of the decision of the patient is necessary for the study and control of the decision of the patient is necessary for the study and control of the decision of the patient is necessary for the study and control of the patient is necessary for the study and control of the patient is necessary for the study and control of the patient is necessary for the study and control of the patient is necessary for the study and control of the patient is necessary for the study and control of the patient is necessary for the study and control of the patient is necessary for the study and control of the patient is necessary for the study and control of the patient is necessary for the study and control of the patient is necessary for the study and control of the patient is necessary for the study and control of the patient is necessary for the study and control of the patient is necessary for the study and control of the patient is necessary for the study and control of the patient is necessary for the study and control of the patient is necessary for the study and control of the patient is necessary for the study and control of the patient is necessary for the study and control of the patient is necessary for the study and control of the patient is necessary for the study and control of the patient is necessary for the study and control of the patient is necessary for the study and control of the patient is necessary for the study and control of the patient is necessary for the study and control of the patient is necessary for the study and control of the patient is necessary for the study and control of the patient is necessary for the study and control of the patient is necessary for the study and control of the patient is necessary for the study and control of t

VIRAL HEPATITIS CASE RECORD FOR REPORTING OF PATIENTS WITH SYMPTOMATIC ACUTE VIRAL HEPATITIS (SEE CASE DEFINITION ON REVERSE)

446800

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STATE	CASE NO).		· · · · · · · · · · · · · · · · · · ·
703		51	(10)	(11)

DEPARTMENT OF HEALTH AND HUMAN SERVICES PUBLIC HEALTH SERVICE

Centers for Disease Control and Prevention Hepatitis Branch, (A33) Atlanta, Georgia 30333

	CDC CASE NO	_
ENTERS FOR DISEASE CONTROL AND PREVENTION	(8) (9	3)

(B) (9) (10) (11)	Atlanta	, Georgia 30333	AND PREVENTION	(8)	(9) (10	<u>)</u> (11)
PATIENT'S LAST NAME (please print clearly) (12-26) FIRST AND	MIDDLE NAME (or initials)	occi	JPATION		
# # # # # # # # # # # # # # # # # # #						
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99 = Unk	9 Unk	ETHINICITY (51) 1 Ha] Unk	
Reporting physician's diagnosis (52-53) 1 DO NOT REPORT CASES OF CHRONIC HER	Hepatitis A 2		(Delta)		patitis ispecified	
CLINICAL DATA			LABORATORY RESUL			
Date of diagnosis (60-65) Was the patient jaundiced? (66) Was the patient hospitalized for hepatitis? (67)	Day Yr / / / / / / / / / / / / / / / / / /	IgM Hepatitis A antibody (Igh Hepatitis B surface antigen (IgM Hepatitis B core antibod Antibody to Della (anti-HDV)	HBsAg) (70) y (IgM anti-HBc) (71)	1	leg Not T	
For purposes of National Surveillance, ASK ALL OF patient acquired his/her infection. Please refer to the	THE FOLLOWING QUES work sheet on the back of	TIONS FOR EVERY CASE OF the last page for additional qu	HEPATITIS. These question uestions.	is may help d	etermine wh	ere the
During the 2-6 weeks prior to illness				Yes	No	Unk
1. was the patient a child or employee in a nursery	, day care center, or presc	hool?		(73) 1 🔲	2 🗌	9 🗌
2. was the patient a household contact of a child of					2 🔲	9 🔲
3. was the patient a contact of a confirmed or susp	ected hepatitis A case? .			(75) 1 🔲	2 🔲	9 🔲
If yes, type of contact: (76) 1 🔲 Sexua	al 2 🔲 Household (non-sexual) 3 🔲 Other				
4. was the patient employed as a food handler?				(77) 1 🔲	2 🗀	9 🔲
5. did the patient eat raw shellfish?				(78) 1 🔲	2 🔲	9 🔲
6. was the patient suspected as being part of a cor	nmon-source foodborne o	r waterborne outbreak?		(79) 1 🔲	2 🔲	9 🔲
7. did the patient travel outside of the U.S. or Cana	ıda?			(80) 1 🔲	2 🔲	9 🗌
If yes, where: (81) 1 So./Central Ar						
		Zealand 7 Other				
Duration of stay: (82) 1 🔲 1-3 Days	2 4-7 Days 3	More than 7 Days				
During the 6 weeks-6 months prior to illness						
8. was the patient a contact of a confirmed or susp	ected acute or chronic he	patitis B or non-A, non-B case?		(83) 1 🔲	2 🔲	9 🔲
If yes, type of contact: (84) 1 Sexua		_				
9. was the patient employed in a medical, dental o	r other field involving conta	act with human blood?		(85) 1 🔲	2 🔲	9 🔲
If yes, degree of blood contact: (86) 1						
10. did the patient receive blood or blood products (-			. (87) 1 🔲	2 🔲	9 🔲
If yes, specify date(s) received: (88-93)						
11. was the patient associated with a dialysis or kid				(100) 1 🔲	2 🔲	9 🗌
		ntact of patient or employee			_	
12. did the patient use needles for injection of stree				.(102) 1 🔲	2 🗌	9 🗌
13. what was the patient's sexual preference? (103					_	
14. how many different sexual partners did the patie				Jnk		
15. did the patient have						
dental work or oral surgery? (105) 1] Yes 2 No	9 ☐ Unk tattooing? .		(108) 1[]	2 🔲	9 🔲
other surgery? (106) 1	_	_ -	tick or puncture with a needle		_	
acupuncture? (107) 1	- 		contaminated with blood? .		2 🔲	9 🔲
Has this patient ever received the three dose ser					2 🔲	9 🔲
If yes, what year? (111-112) A	ND was the nationt tester	for antibody within 1-6 months	after the last dose?	(113) 1	2	9 🔲
If yes, was the antibody test: (114) 1		3 Unknown		. ,	_	
Comments:			Investigator's Name	<u> </u>		
			Date			

WORK SHEET

CASE DEFINITION FOR REPORTING OF ACUTE VIRAL HEPATITIS

Illness with: 1) discrete onset of symptoms and

2) jaundice or elevated serum aminotransferase levels.

Hepatitis A: IgM anti-HAV positive.

Hepatitis B: IgM anti-HBc positive if done or HBsAg positive and IgM anti-HAV negative if done

Non-A. Non-B Hepatitis: 1) IgM anti-HAV negative, and

2) IgM anti-HBc negative if done or HBsAg negative, and

3) serum aminotransferase levels greater than 2 1/2 times the upper limit of normal

<u>Delta Hepatitis:</u>

1) HBsAg or IgM anti-HBc positive and

2) Anti-HDV positive.

FOR USE BY LOCAL HEALTH	DEPARTMENTS TO DETER	MINE THE PATIENTS M	OST PROBABLE SO	OURCE OF INFECTION
Patient's name	Home phone	Employed by		Work phone
Reporting physician's name, address, and	phone #			
If patient was hospitalized for hepatitis, given				
Results of liver function tests: SGOT (AS	ST)SGP	T (ALT)	Bilirubin	
	MATION FOR ADMITTED RISI			ONT PAGE
	MATION FOR ADMITTED RISI	C PACTORS AND SOON	020 2.0 (20 0)	
IF APPLICABLE: 1. Name, address, and phone # of c	child care center			
مع المعظمة في مستقلات الناب الناب الناب	do classroom attended			
Name and address of school, gra Name, address, and phone # of re	estaurant where food handler v	vorked (HEPATITIS A O	NLY)	
4. Food history of patient for the 2-6	wks prior to onset: (HEPATIT	IS A ONLY)		
a. name and location of restaur	ants			
 name and location of food st 	ores			
c. name and location of bakery	i lumb marting ato	\		
d. group meals attended (e.g., r	sed	.)		
e. location raw shellfish purcha5. Name, address, and phone # of least to the properties of the purchase.	sed	R contact		
Name, address, and phone # of F	known nepatitis A of Nepatitis E	o comaci	Relationship	
CONTACTS	REQUIRING PROPHYLAXIS F	OR HEPATITIS A OR HE	PATITIS B	
6. CONTACTS I		ship to case IG	HBIG	Vaccine
Wallio .			·	
7. If transfused, NOTIFY BLOOD C	ENTER! Name of blood cer	nter		
a. number of units of whole blo	od, packed RBC or frozen RB0	C received		
b. specify type of blood produc	t (e.g., albumin, fibrinogen, fac	tor VIII, etc.)		
8. IF DONOR, name, address, and	phone # of donor or plasmapi	neresis center		Date
9. Name, address, and phone # of	dialysis center			
10. Name, address, and phone # of	dentist or oral surgeon	tion		
11. If other surgery performed, name	e, address, and phone # of loca	ation		
12. Name, address, and phone # of	acupulacturist or tatlog parter			
12. Name, address, and phone # of13. Is patient currently pregnant?	acupunctions: or tation partor	rian's name, address and	phone #	
13 Is patient currently pregnant?	ii yes, give obstetii	San S name, accress and		
a. estimated date and location	of delivery			
 a. estimated date and location 	ur denvery			
Comments:				
Comments.				
Investi	gator's Name and Title			Date of Interview

Work sheet